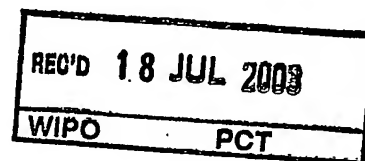




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10 Nov 03 PCT/PTC 03 JAN 2005



Patent Office
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I, JULIE BILLINGSLEY, TEAM LEADER EXAMINATION SUPPORT AND SALES hereby certify that annexed is a true copy of the Provisional specification in connection with Application No. PS 3352 for a patent by USCOM PTY LTD as filed on 03 July 2002.



WITNESS my hand this
Seventh day of July 2003

JULIE BILLINGSLEY
TEAM LEADER EXAMINATION
SUPPORT AND SALES

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AUSTRALIA

PATENTS ACT 1990

PROVISIONAL SPECIFICATION

FOR THE INVENTION ENTITLED:-

"Prosthetic Heart Function Evaluation Method and Apparatus"

The invention is described in the following statement:-

Prosthetic Heart Function Evaluation Method and Apparatus

Field of the Invention

The present invention relates to the field of monitoring of prosthetic heart devices and in particular, disclose the form of a CW Doppler monitoring of such
5 devices.

Background of the Invention

The malfunctioning of the heart muscle is a major killer in society. Various devices have been proposed for assisting the operations of the cardiac function and stimulating the heart's natural pumping action. These heart assist devices
10 come in many forms from full heart replacement to minor forms of assist, where small pumps connected to the ventricle are mounted outside the heart, and connecting flow from the ventricle to the aorta, by-passing the ventricle and systemic aortic valve.

In Fig. 1, there is shown a perspective view partly in section of the human
15 heart. The heart can be divided into four chambers including the right atrium 2, the right ventricle 3, the left atrium 4 and the left ventricle 5. Between the chambers 2, 3 is the tricuspid valve 6. At the exit to the right ventricle is the pulmonary valve 7. Between the two chambers 4, 5 is the mitral valve 8. At the exit to the left ventricle is the aortic valve 9.

20 Although many different prosthetic devices exist, including complete heart replacement, one form of prosthetic device bypasses the native left ventricular circulation by redirecting flow from the left ventricle 5 to the thoracic aorta 11. These prosthetic devices are designed to offload the ventricle 5 by redirecting flow from the ventricle mechanically, and encouraging the native ventricle to
25 increase its function under decreased load.

It would be desirable to provide for accurate flow monitoring of such a system for the analysis of the proper functioning of such devices.

Summary of the Invention

It is an object of the present invention to provide for an improved form of
5 monitoring of the flow through the heart in conjunction with a prosthetic assist device.

In accordance with a first aspect of the present invention, there is provided a method of monitoring the operation of a prosthetic assist device, the method comprising the steps of: (a) utilising a non-invasive device to monitor directly the
10 blood flow through the by-passed heart ventricle; (b) separately monitoring the blood flow through the prosthetic assist device; (c) combining said two measurements to determine an overall cardiac output and a native to prosthetic flow ratio.

Preferably, the non-invasive device monitoring comprises continuous wave
15 Doppler flow monitoring of the heart. The monitoring can occur from a transducer placed over the parasternal acoustic access and can be repeated under a number of different operational conditions for a patient including walking and/or running. Ideally, the method is repeated under a number of different pharmacological conditions for a patient.

20 Brief Description of the Drawings

The preferred embodiments of the present invention will now be described with reference to the accompanying drawings in which:

Fig. 1 is a perspective view partly in section of the human heart;

Fig. 2 illustrates schematically the operation of a prosthetic assist device;

Fig. 3 illustrates utilisation of a CW transducer device for monitoring heart function;

Fig. 4 illustrates the transducer device of Fig. 3;

Fig. 5 illustrates one form of output of the transducer device; and

5 Fig. 6 illustrates the various portions output in the arrangement of Fig. 5.

Description of the Preferred and Other Embodiments

In the preferred embodiment, accurate measurement of total ventricular function is provided by continuous wave Doppler (CW Doppler) measurement of the native ventricular function in addition to accurate flow measurement through
10 the mechanical device.

Turning initially to Fig. 2, there is illustrated a schematic of the human heart
10 which includes the aorta 11 and an additional mechanical assist device 12 which has a shunted ventricle input 13 and an aorta output 14. Whilst the flow through the mechanical assist 12 can be accurately measured, the flow out of
15 the heart 10 via aorta 11 is less accurately measurable. In the preferred embodiment, techniques of CW measurement are provide so as to provide for flow measurements produced by the heart.

Turning to Fig. 3, there is shown a patient 20 having a non-invasive sensor 21 attached to the body. The sensor 21 comprises a CW transducer device
20 which is interconnected to a base station unit 22.

Fig. 4 shows an example of the first sensor transducer actuator 11 for attachment to the skin surface. CW Doppler measurements are utilised to monitor the blood flow through the heart. CW Doppler is a non-invasive technique in which ultrasonic signals from transducer elements are directed into
25 a blood carrying vessel of a patient. Doppler shifts in the reflected signal

provide an indication of the rate of blood flow. In Fig. 2, a transducer element 21 includes an ultrasonic transducer 25 attached to a positioning device 26 which can be used to initially set the position of the transducer. Between the transducer 25 and a patient's skin 27 is placed a gel coupling layer 28 for
5 coupling the ultrasonic transducer vibrations to the skin 27. The principles of CW Doppler flow measurement are known and do not themselves form part of the present invention. Patent Cooperation Treaty (PCT) publication number WO 99/66835 entitled "Ultrasonic Cardiac Output Monitor" assigned to the present assignee, the contents of which are incorporated herein by cross-reference,
10 describes in more detail an ultrasonic transducer device suitable for measuring blood flow within the heart using the CW Doppler method. In the embodiment shown in Fig. 3, the transducer elements are placed on the patient to obtain intra-cardiac or aortic signals, for example through a suprasternal notch.

The CW method detects the velocity of individual blood cells by measuring
15 the frequency change of a reflected ultrasound beam and displaying this as a time velocity flow profile, an example of which is shown in Fig. 5. The transducer output forms an input to the processor unit 22 of Fig. 3. From the time velocity flow profile, the processor can calculate the time velocity integral (tvi) and other relevant information such as heart rate (HR), each of which is
20 illustrated in Fig. 6. The information can be derived from Fig. 6 by appropriate image processing of the CW output image.

The total systemic output, or the total volume of blood directed into the systemic vessels, will be the sum of the native plus the prosthetic output. The ratio of the two flows can be denoted a native to prosthetic flow index and can

be utilised to determine appropriate levels for setting the prosthetic output of the prosthetic device 12 of Fig. 2.

Preferably, after fitting the heart assist device, the performance of the device relative to heart rate measurements are conducted under a number of
5 different conditions.

Through measuring the device performance for a wide range of patients under a wide range of different conditions, an extensive database can be constructed of optimal performance conditions. In this way, the heart assist device can be tuned for sophisticated operation.

10 Further, the utilisation of the index and extensive measurements allows for optimisation of pharmacotherapy. The monitoring device thereby provides the ability to accurately monitor output flow from the heart function with potential therapeutic benefits.

It will be understood that the invention disclosed and defined herein
15 extends to all alternative combinations of two or more of the individual features mentioned or evident from the text or drawings. All of these different combinations constitute various alternative aspects of the invention.

The foregoing describes the embodiments of the present invention and modifications, obvious to those skilled in the art can be made thereto, without
20 departing from the scope of the present invention.

The claims defining the invention are as follows:

1. A method of monitoring the operation of a prosthetic assist device, the method comprising the steps of:

5 (a) utilising a non invasive device to monitor directly the blood flow through at least one heart ventricle;

(b) separately monitoring the blood flow through the prosthetic assist device;

(c) combining said two measurements to determine an overall native to prosthetic flow index.

2. A method as claimed in claim 1 wherein said non invasive device
10 monitoring comprises continuous wave Doppler flow monitoring of the heart.

3. A method as claimed in claim 1 wherein the heart is monitored from a transducer placed adjacent the suprasternal notch.

4. A method as claimed in claim 1 wherein said method is repeated under a number of different operational conditions for a patient including walking and/or
15 running.

5. A method as claimed in claim 1 wherein said method is repeated under a number of different pharmacological conditions for a patient.

DATED this 3rd day of July, 2002

USCOM PTY LTD

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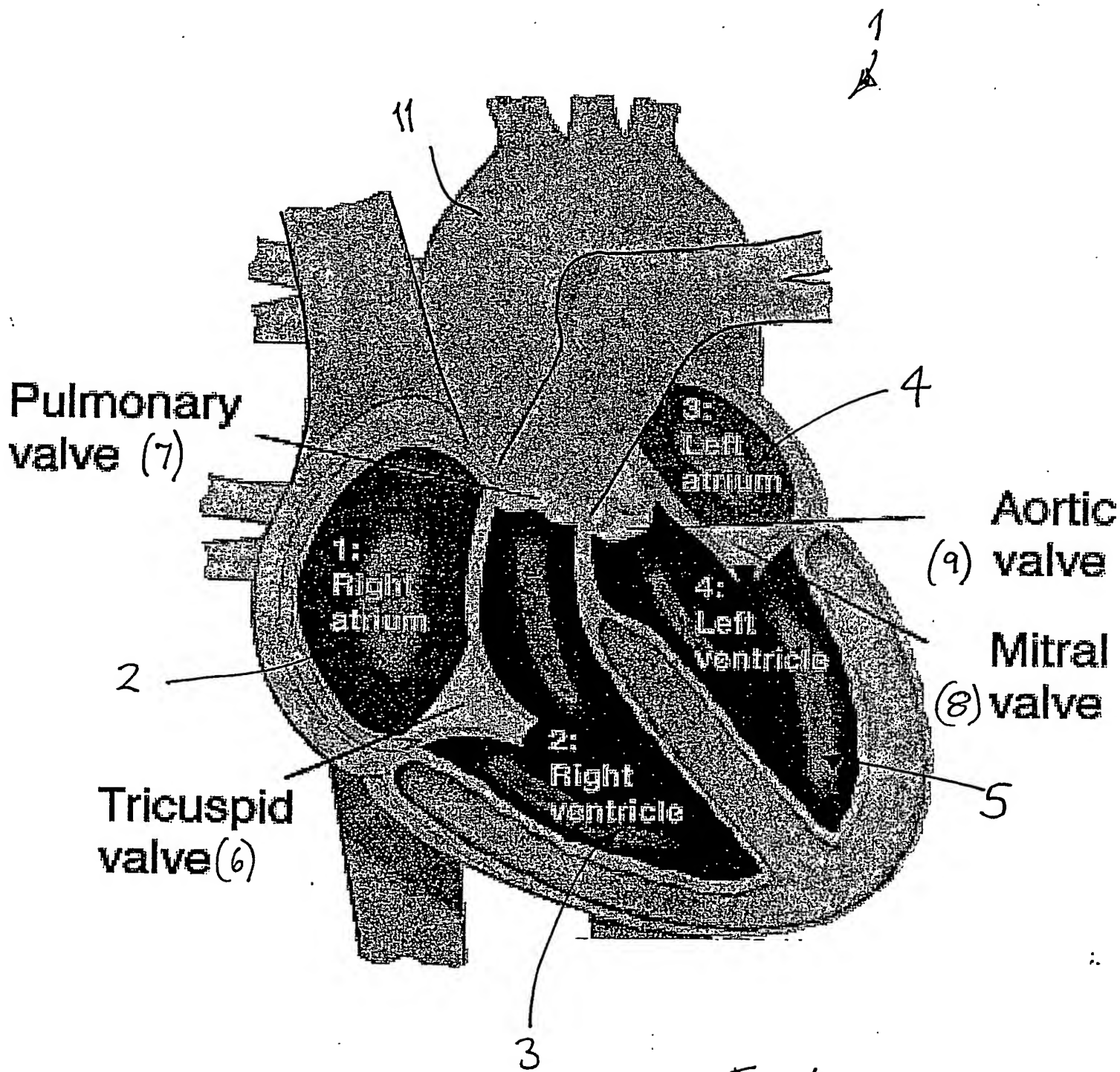


Fig. 1

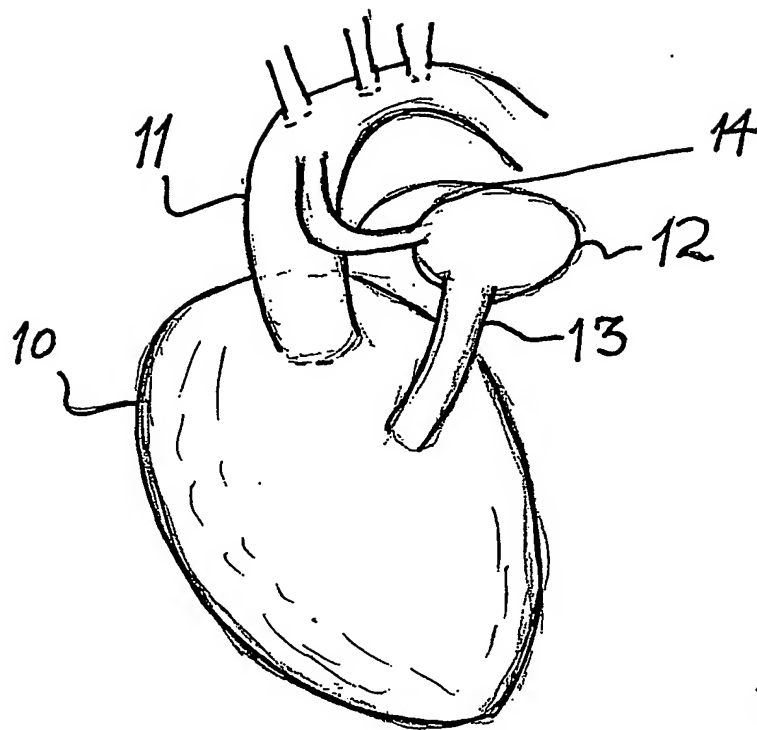


FIG. 2

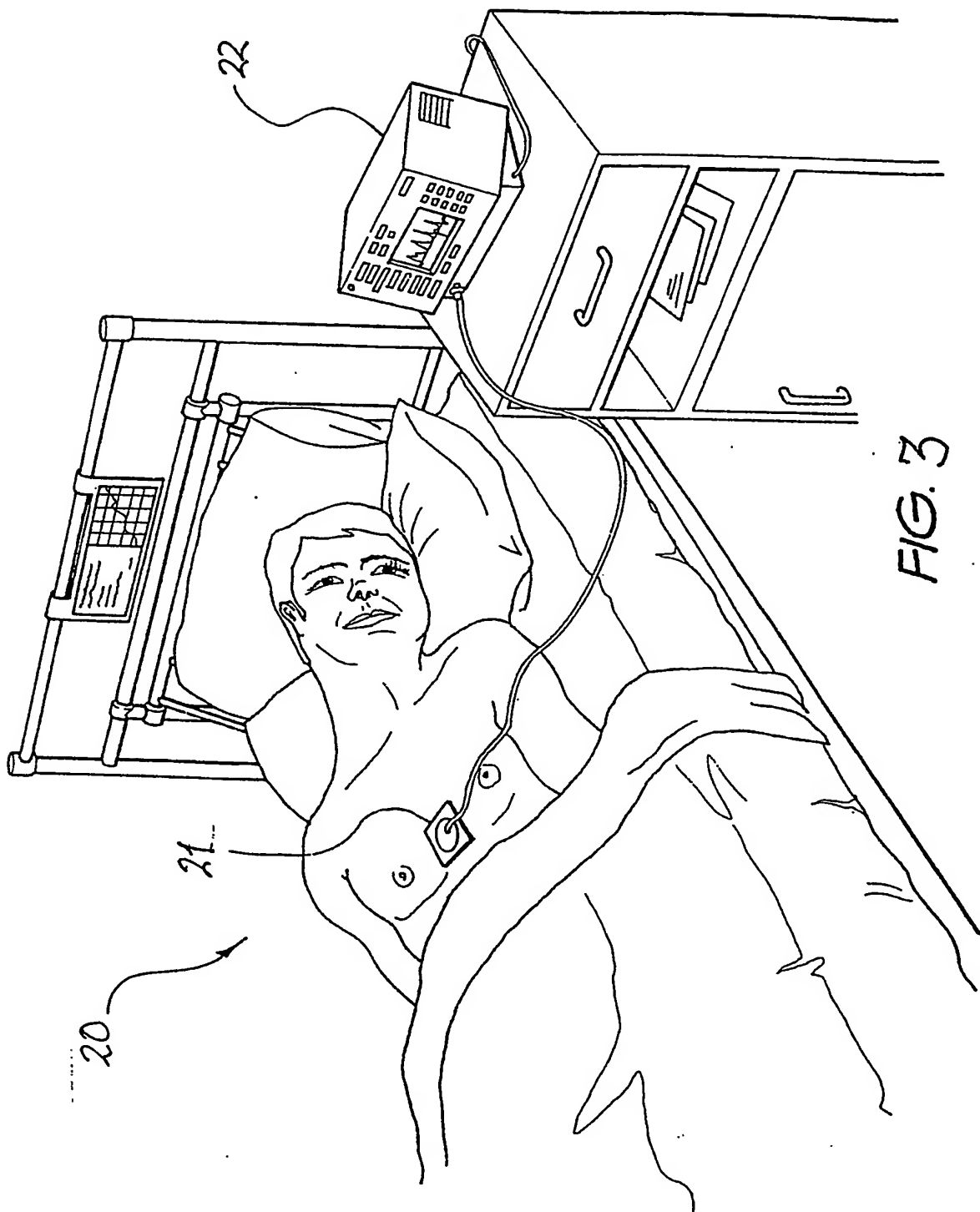


FIG. 3

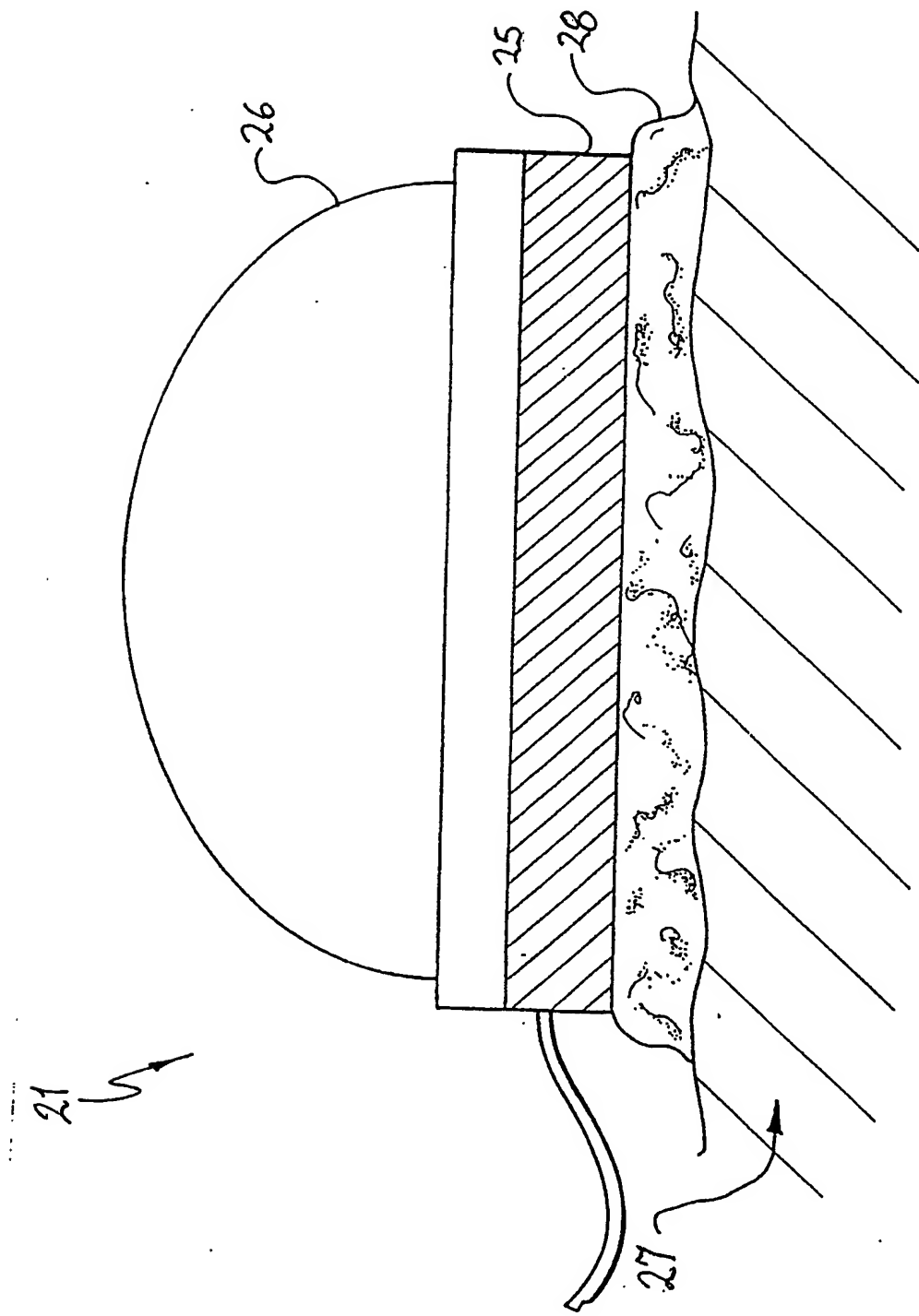


FIG. 4

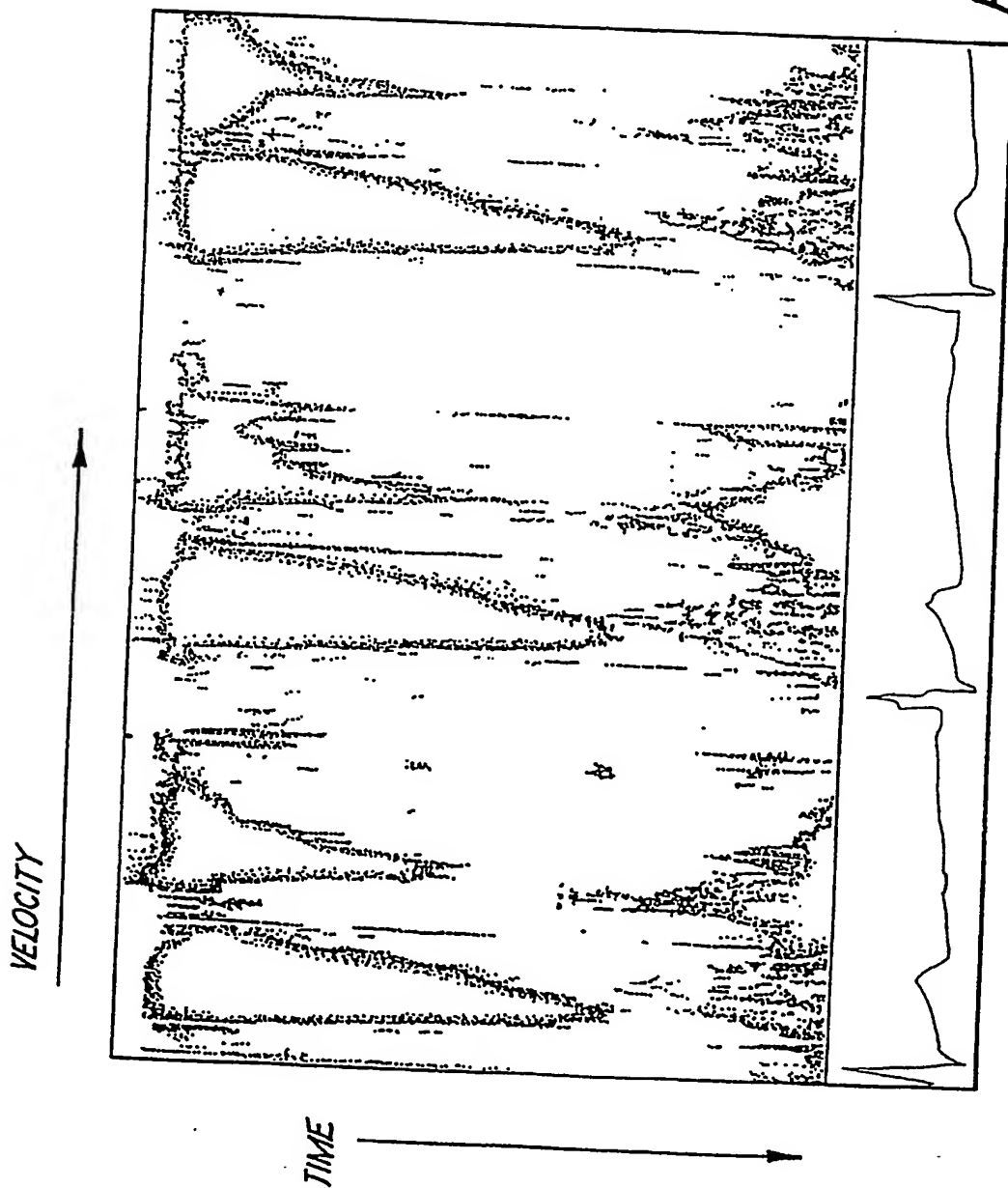


FIG. 5

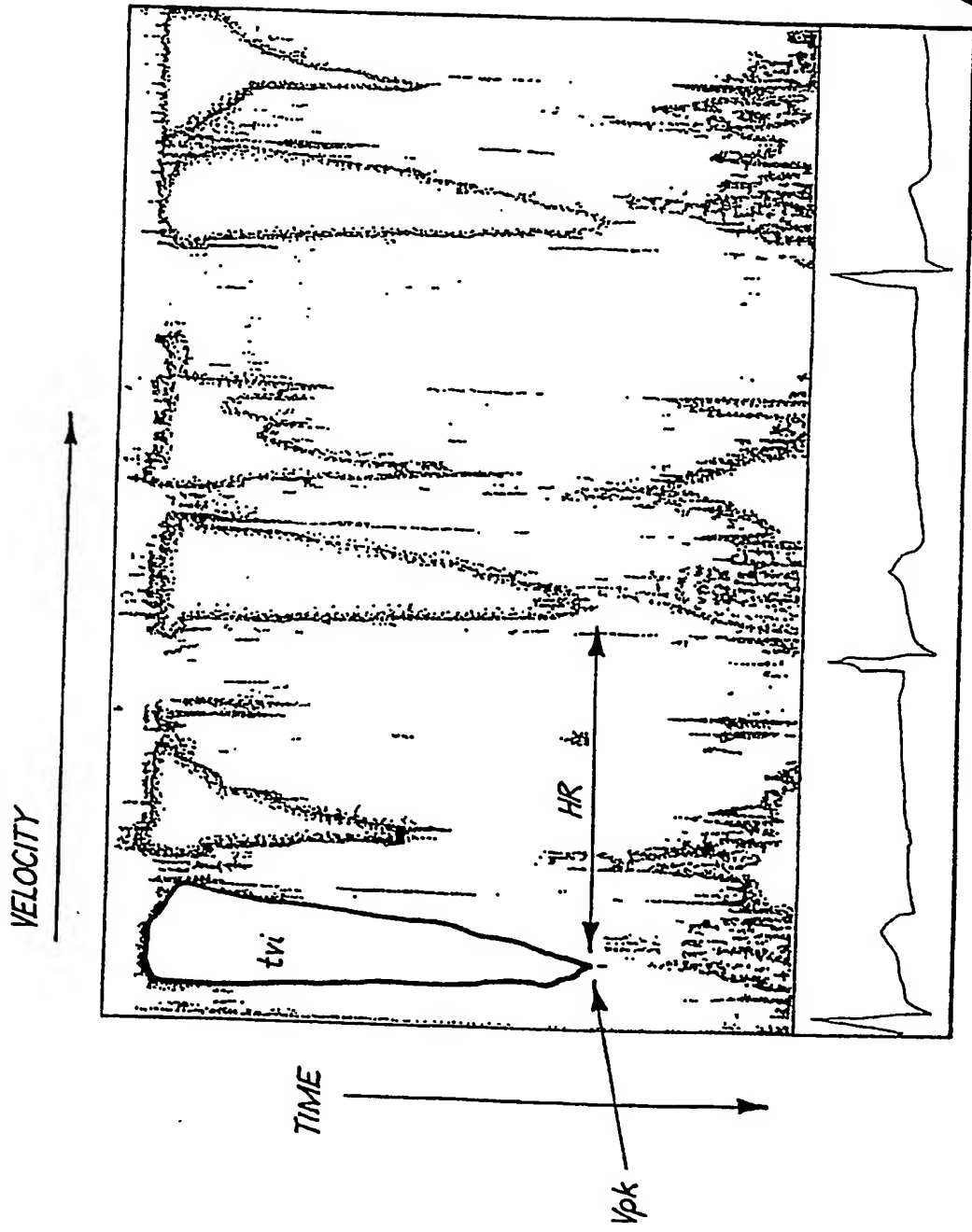


FIG. 6